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Mechanical properties and deformation behavior of cast binary Ti-Cr alloys

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1* Identify the short form here as you did in the 2nd sentence of the Introduction: "commercially pure titanium."

Abstract

As-cast Ti-Cr alloys prepared using a dental cast machine with Cr contents ranging from 5 [space?] were Obviously to 30 wt% have been investigated. Experimental results indicated that the Ti-Cr alloys had [redundant] obviously different deformation behavior with various amounts of Cr contents. The bending strength of the Ti-20Cr alloy was about 1.8 times greater than for c.p. Til Ti-10Cr alloy had * 1 as much as relatively higher bending strength almost near the Ti-20Cr alloy. This is believed to be a 2* result of the strengthening effect of ω phase. From SEM fractographs, the Ti-10Cr alloy was Marasterized featured by coarse cleavage facets in the fracture surface together with some terrace-type colony howevery comma morphology. The Ti-20Cr exhibited ductile properties, but not for alloys with other did not exhibit such properties. compositions. In addition, the elastic recovery capability of Ti-20Cr alloy was greater than that of c.p. Ti by as much as 460%. From the unetched optical micrographs, the surfaces of (commatherebyin numbers Ti-20Cr alloys were covered with large amounts of slip bands, It showed that the deformation of Ti-20Cr alloy was dominated by slip of dislocations. By judging from the results of mechanical properties and deformation behavior, Ti-20Cr is considered to be the most Cosible expected alloy for prosthetic dental applications if other properties necessary for dental casting are obtained.

Keywords: Titanium alloy; Chromium; & phase; Mechanical properties

2* Identify the acronym "SEM": scanning electron microscopy (according to p.5).

3* Use "number(s)" with countable nouns such as "bands" / "amount(s)" with uncountable nouns such as "content."

1* Should the syntax be % wt throughout this report? The phrase "wt %" does not sound correct to me.

1. Introduction

Titanium and titanium alloys have become one of the most attractive biomaterials due to their light weight, high biocorrosion resistance, biocompatibility and mechanical properties.

Commercially pure titanium (c.p. Ti) has also established a reputation for excellent biocompatibility as a dental metal and is suitable for dental prosthetic applications [1,2,3].

There are some inherent problems that must be overcome for titanium to be used successfully in dentistry. The fact that pure titanium has a high melting temperature and high reactivity with oxygen and impurities at elevated temperatures makes it difficult to cast [4]. In addition, reducing the melting temperature of titanium also could decrease the risk of inadequate mold filling and porosity development due to the considerable temperature difference between the molten alloy and the much cooler investment [5]. On the other hand, since the properties of unalloyed titanium are not suitable for all purposes, many titanium alloys for dental use have been developed, and their properties have been studied [1,6-9], mainly to improve the strength and castability of pure titanium.

One method of solving these problems is to use titanium alloys, which could exhibit solid solution hardening and have lower fusion temperatures and better ductility than c.p. Ti [6].

Titanium can also be alloyed with a variety of elements to alter its properties, namely, to improve strength, high temperature performance, creep resistance, weldability and formability [1]. For a metal to be used in a dental restoration, it should be biocompatible so that it does not cause harmful toxicological or allergic reactions to the patient. Among the various titanium alloys, Ti-6Al-4V alloy is the most widely used because of its better physical and mechanical properties in comparison to c.p. Ti [1]. However, there has been a speculation that the release of Al and V ions from the alloy might cause some long-term health problems [10-12]. Moreover, the low wear resistance of Ti-6Al-4V could accelerate the release of such harmful ions [13-15]. The searches for alternatives capable of satisfactorily replacing the

1* "Research" is uncountable. Do not attempt to make it plural! Furthermore, uncountable nouns cannot take an indefinite article.

traditionally used alloys made titanium become a target for researches in prosthetic dentistry.

In the 1980s, Ti-Cr alloys were reported as having high tensile strength and good ductility for dental casting alloys as well as Ni-Cr and Co-Cr alloys [16,17]. In our previous study [18], the hardness of a series of binary Ti-Cr alloys with Cr contents ranging from 5 to 30 wt% had was been investigated. We found that the hardness of the Ti-Cr alloys became higher as the Cr contents increased, and ranged from 296 HV to 428 HV. The present study is a continual research of this alloy system, with the focus on the mechanical properties and the deformation behavior of these titanium alloys. Also, the mechanical properties of the cast Ti-Cr alloys were investigated to judge the possibility for the practical dental applications.

2. Materials and Methods

The materials used for this study include c.p. Ti, Ti-5Cr, Ti-10Cr, Ti-20Cr, Ti-25Cr and Ti-30Cr alloys (in wt%). Based on the Ti-Cr phase diagram [19], the eutectoid composition is approximately at 13.5 wt%. The Ti-5Cr and Ti-10Cr alloys correspond to the hypoeutectoid compositions, and Ti-20Cr, Ti-25Cr and Ti-30Cr alloys are in the hypereutectoid composition range. All the materials have been prepared from raw titanium (99.8% in-purity), Cr (99.95% in-purity) using a commercial arc-melting vacuum-pressure-type casting system (Castmatic, Iwatani Corp., Japan). The ingots of approximately 20 g each were re-melted five times to improve chemical homogeneity. Prior to casting, the ingots were re-melted again. The difference in pressure between the two chambers allowed the molten alloys to instantly drop into a graphite mold at room temperature. In this study, the experimental alloys were fabricated using a graphite mold instead of dental investment mold.

X-ray diffraction (XRD) for phase analysis was conducted using a diffractometer (XRD-6000, Shimadzu, Japan) operated at 30 kV and 30 mA. A Ni-filtered CuKa radiation was used for this 2* wherein the phases were study, Phase was identified by matching each characteristic peak with the JCPDS files. 3* 2* Radiation is an uncountable noun. Do not precede with an indefinite article. 3* Where is your identification of the across "JCPDS"?

*Is this the typical method for calculating the modulus of elasticity in bending? If so, "is" is correct. However, if this method can typically vary with the experiment, then "was" should be used.

Three-point bending tests were performed using a desk-top mechanical tester (AG-IS, Shimadzu, Japan). The bending strengths were determined using the equation, $\sigma = 3PL/2bh^2$ [20], where σ is the bending strength (MPa), P is the load (kg), L is the span length (mm), b is the specimen width (mm), and h is the specimen thickness (mm). The dimensions of the specimens were L = 30 mm, b = 5.0 mm and h = 1.0 mm. The modulus of elasticity in bending is calculated from the load increment and the corresponding deflection increment between the two points on the straight line as far apart as possible using the equation E = $L^{3}\Delta P$ /4bh³ $\Delta\delta$, where E is the modulus of elasticity in bending (Pa), ΔP is the load increment as measured from preload (N), and $\Delta\delta$ is the deflection increment at midspan as measured from preload. The average bending strength and modulus of elasticity in bending were taken Sive from at least 5 tests under each condition. The elastic recovery (springback) capability for each material was evaluated from the change in deflection angle when loading was removed. Details can be found in Ho et al. [21]. The fracture surface of the fractured specimen after the bending test was cleaned by an ultra sonic washer. The fracture surface was observed using s scanning electron microscopy (SEM; JSM-6700F, Jeol, Tokyo, Japan). In addition, microstructures of post-bending unetched specimens were examined using an optical

3. Results and discussion

microscope (BH2, Olympus, Japan).

3.1. Phase and structure

Phase and crystal structure of c.p. Ti and Ti-Cr alloys was shown is Table 1. The crystalline structure of the binary Ti-Cr alloy is sensitive to the composition (chromium content) of the alloy. The c.p. Ti was comprised entirely of a hexagonal α phase. With 5 wt% Cr, metastable β phase starts to be retained. When the Cr content increased to 10 wt% or higher, the β phase was completely retained with a bcc crystal structure. This finding is consistent with the early what is "bcc"?

(at"?

result of Ericsen et al. [22]. They reported that the β phase cannot be fully retained during quenching unless the Cr content exceeds about 6 at% (6.5 wt%). Apparently, more than 10%Cr is needed to fully retain the β phase in the Ti-Cr alloys. Like Fe, Co and Ni, Cr has been recognized as an eutectoid β-stabilizing element. This is true for the present Ti-Cr system.

The ω peak was found in the Ti-5Cr and Ti-10Cr alloys, especially notable in Ti-10Cr alloy. According to Sikka et al. [23], this ω phase may be defined by a hexagonal lattice. In many investigations, the presence and volume fraction of an ω phase in alloys of titanium and zirconium can be seen from the intensity of lines on the x-ray diffraction patterns of polycrystals or of one of the ω phase reflections in photographs of single crystals [24-26]. The presence of this athermal ω phase, though being small in quantity, had an exceedingly important effect on the mechanical properties of the alloy, as will be discussed later.

3.2. Mechanical properties

The bending strengths of c.p. Ti and Ti-Cr alloys are shown in Fig. 1. ANOVA test results showed significant overall differences among bending strengths for c.p. Ti and Ti-Cr alloys (p < 0.05). All the Ti-Cr alloys had significantly higher (p < 0.05) bending strength (1035-1484 MPa) than the c.p. Ti (844 MPa) tested. The bending strengths of the Ti-Cr alloys showed the highest results in the alloy having 20 wt% Cr content. The bending strength of the Ti-20Cr alloy was about 1.8 times greater than for c.p. Ti. Ti-10Cr alloy had relatively higher bending strength almost near the Ti-20Cr alloy. This is believed to be a result of the strengthening effect of ω phase. The ω phase occurs in some titanium base alloys in which β phase can be retained in a metastable state [25].

The elastic modulus results are shown in Fig. 2. ANOVA test results showed significant overall differences among bending modulus values for c.p. Ti and Ti-Cr alloys (p < 0.05). For

the Ti-Cr alloys, the elastic modulus values had a tendency to increase as the chromium content increased. It is worth noting that the elastic modulus of Ti-10Cr alloy (163 GPa) was significantly higher (p < 0.05) than all the other Ti-Cr alloys and c.p. Ti. This result may be concerned with the formation of the ω phase during quenching. The early work of Graft et al. However [27] indicated that the ω phase has an unusually high elastic modulus. On the other hand, although the bending strength seems to slightly decrease with Cr content for higher Cr alloys, their moduli increased significantly. For example, when Cr content increased to 30 wt%, the modulus of the alloy increased to 147 GPa. These results indicate that, at high Cr contents, modulus is more sensitive to Cr content than strength. The similar results had also been discussed on the Ti-7.5Mo-xFe alloys [28].

Typical bending stress-deflection profiles of the series of alloys are shown in Fig. 3. White Ti-5Cr and Ti-10Cr alloys with ω phase failed in a brittle way (with an average deflection of only about 2.4 and 1.8, respectively), Ti-20Cr alloy without ω phase did not fail even after being deflected by 8 mm (the pre-set maximum). This ω phase-induced embrittlement has also been found in other Ti alloy systems as early as 1970s, such as Ti-V [29] and Ti-Mn [30]. Recently, Lin et al. also reported the mechanical behavior of the ω phase in Ti-7.5Mo-xFe alloy system [31]. It is interesting to note that, despite the strong hardening effect of w phase, Ti-locry commas the bending strength of the alloy comprising the largest amount of ω phase (Ti-10Cr) was slightly lower than those containing no ω. This is due to the premature, brittle fracture that occurred for Ti-10Cr. The embrittling/weakening effect was very sensitive to the composition, or to be exact, to the Cr content of the alloy. As shown in Fig. 3, when the Cr content was space] higher or lower than 20 wt%, the alloys showed brittle properties. In this series of alloys the space? highest bending strength was found in the alloy around 20 wt%Cr (1484 MPa). Aperiod an abrupt decrease in stress

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* Avoid the phrase "and then" in Sormal writing. Both words are connectors. Only one word is needed. Do not overwork the word "and." 7

* deflection of about 2 mm. And then, the stress gradually decreased, which could be related to

The stress-deflection curve of Ti-25Cr alloy exhibited a stress decrease abruptly at the a

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bending strength and deflection were found for the Ti-30Cr alloy. For the Ti-25Cr and the Ti-30Cr alloys, the fractureness is believed to be the result of the inclusion of the eutectoid consistuents. TiCr₂ and/or Ti₂Cr₃ in the microstructure. However, the XRD failed to find these constituents, probably due to the limited detection capability of the x-ray diffractometer. This finding concurs with those of Koike et al. [32].

In this study, only

The

The Ti-20Cr exhibited ductile properties, but not for alloys with other compositions in this study. As indicated in the results of Koike et al. [32], increasing Cr to 19% produced the best ductility in the alloys by tensile testing. The tensile test data for binary Ti-Cr alloys found earlier by Okuno et al. [6] were 764 MPa (Ti-10Cr) and 970 MPa (Ti-20Cr); the elongation values were 0.2% (Ti-10Cr) and 2.8% (Ti-20Cr). It is worth noting that the advantage in mechanical properties of Ti-20Cr alloy is also demonstrated in their high elastic recovery capability. High elastic recovery (springback) capability of a metal is an indication of high strength and low modulus and is essential for many load-bearing implant and dental applications. The elastic recovery capability of Ti-20Cr alloy was greater than that of c.p. Ti by as much as 460%.

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3.3. SEM photography

Fig. 4(a)-(d) shows SEM micrographs of the fractured surfaces of the Ti-Cr specimens after bending test. Since Ti-20Cr alloy did not fail during bending test, its micrograph was not examined. The Ti-10Cr alloy was featured by coarse cleavage facets in the fracture surface, which are characteristic of decreased ductility, together with some terrace-type morphology. At higher magnification (Fig. 5(a)), the terrace-type morphology was more obvious, similar to that observed in a recent report of Lin et al. [31]. The fractured structures of the Ti-25Cr alloy exhibited

(Fig. 4(c)) mainly cleaved grains. On the other hand, dimple ruptures, indicative of a typical

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ductile fracture, were observed in the fractured structures of the Ti-25Cr alloy at a higher magnification view of SEM fractographs (Fig. 5(b)). Also, the fractured surface showed some dimple ruptures with some coalescence of microvoids. The fractured surface morphology of the Ti-30Cr alloy (Fig. 4(d)) revealed completely cleaved grains. The cleavage fracture is in corresponds to agreement with the highly brittle failure of this specimen indicated by the extremely low value of bending deflection (roughly about 1.5 mm). As shown in Fig. 4(a), the fractured surface topography of Ti-5Cr alloy revealed cleaved grains with some coalescence of sub-micron-sized dimples, and the cleaved grains were much smaller than that of Ti-10Cr or the Ti-30Cr alloy. The cleavage nature of the fracture seen in these fractographs is consistent with being that of the a fracture deflection of 2.4 mm exhibited by this specimen, which are larger than Ti-10Cr (1.8 mm) and Ti-30Cr (1.5 mm) alloys.

3.4. Optical micrographs of as-bent surfaces

Fig. 6 compared the as-bent convex side (tensile stress side) surface morphologies of the indicate.

Ti-Cr alloys. It can be seen from these unetched optical micrographs that the surfaces of the Ti-5Cr, Ti-20Cr and Ti-25Cr alloys were covered with large amounts of slip bands, especially obvious for the Ti-20Cr alloy. Malso showed that the deformation of Ti-20Cr and Ti-25Cr alloys was dominated by slip of dislocations. In addition, Ti-25Cr alloy showed the cracks the along the grain boundaries. On the other hand, the surfaces of Ti-10Cr and Ti-30Cr alloys did reveal the tension of the surfaces of the tension of the plastic deformation.

For a metal to be used as a clasp on removable prosthetics, high elongation is desirable to minimize the risk of breakage. Also, if such an alloy is accidentally stressed beyond its proportional limit, a high elongation value will insure against fracture [33]. Alloys exhibiting high elongation values coupled with high tensile strengths usually have good fracture

* In this context, "literature" is an uncountable noun. Do not attempt to make it plural

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resistance [34]. Thus, we should not use brittle alloys for dental applications even if their strength is sufficient. Therefore, from an engineering point of view, not only a Cr content of 5, 10, 25 and 30 wt% should be avoided, but also a uniform distribution in Cr is also important for such an alloy system. When any process-induced segregation occurred, ω-induced embrittlement may occur in any local regions which happen to have a Cr concentration away from 20 wt%.

Our study indicated that the Ti-20Cr alloy did not exhibit the feature of breaking or cracking the during bending test before the deflection of 8 mm, which may reduce the risk of breakage or fracture of partial dentures in clinical use. According to the literatures, Oda et al. [35,36] reported that an experimental cast of Ti-20Cr alloy had the least tarnish when immersed in an moreover acidic saline solution containing fluoride or hydrogen peroxide. [Also], Takemoto et al. [37] found stated that the Ti-20Cr alloy had a greater resistance to corrosion in a fluoride-containing saline solution than did c.p. Ti due to formation of a chromium-rich oxide film. Consequently, when by judging from the results of mechanical properties and deformation behavior in this study, Feasible

Ti-20Cr alloy is considered to be the most expected alloy for prosthetic dental applications if other properties necessary for dental casting are obtained.

4. Conclusions

- (1) The bending strengths of all the Ti-Cr alloys (1035-1484 MPa) were significantly higher than that of c.p. Ti (844 MPa). The bending strength of the Ti-20Cr alloy was about 1.8 that of times greater than for c.p. Ti. Ti-10Cr alloy had relatively higher bending strength almost that of near the Ti-20Cr alloy. This is believed to be a result of the strengthening effect of the phase.
- (2) The elastic modulus values of the Ti-Cr alloys had a tendency to increase as the chromium content increased. The elastic modulus of Ti-10Cr alloy (163 GPa) was significantly

higher than all the other Ti-Cr alloys and c.p. Ti. This result may be concerned with the formation of the ω phase during quenching.

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the study. The elastic recovery capability of Ti-20Cr alloy was greater than that of c.p. Ti by

as much as 460%.
After a bending test images

(4) SEM micrographs of the fractured surfaces after bending test showed that the Ti-10Cr alloy was featured by coarse cleavage facets in the fracture surface, which are characteristic of decreased ductility, together with some terrace-type morphology.

(5) From the unetched optical micrographs, the surfaces of Ti-20Cr alloys were covered with numbers large amounts of slip bands. It showed that the deformation of Ti-20Cr alloy was dominated by slip of dislocations.

Acknowledgment

Department of English, Da Yeh University,

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Table and figure captions

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Table 1 Phase and crystal structure of c.p. Ti and Ti-Cr alloys fabricated in this study.

- Fig. 1. Bending strengths of c.p. Ti and Ti-Cr alloys.
- Fig. 2. Bending moduli of c.p. Ti and Ti-Cr alloys.
- Fig. 3. Bending stress-deflection profiles of c.p. Ti and Ti-Cr alloys.
- Fig. 4. SEM fractographs of Ti-5Cr (a), Ti-10Cr (b), Ti-25Cr (c) and Ti-30Cr (d) alloys.
- Fig. 5. SEM fractographs of Ti-10Cr (a), Ti-25Cr (b) alloys.
- Fig. 6. Optical micrographs of as-bent surfaces of Ti-5Cr (a), Ti-10Cr (b), Ti-25Cr (c) and Ti-30Cr (d) alloys.

Table 1 Phase and crystal structure of c.p. Ti and Ti-Cr alloys fabricated in this study.

Alloy	Phase	Crystal structure
c.p. Ti	α	hexagonal
Ti-5Cr	α/β/ω	hexagonal / bcc / hexagonal
Ti-10Cr	β/ω	bcc / hexagonal
Ti-20Cr	β	bcc
Ti-25Cr	β	bcc
Ti-30Cr	β	bcc

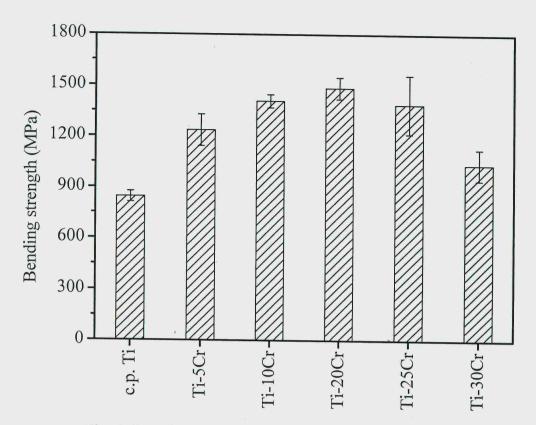


Fig. 1. Bending strengths of c.p. Ti and Ti-Cr alloys.

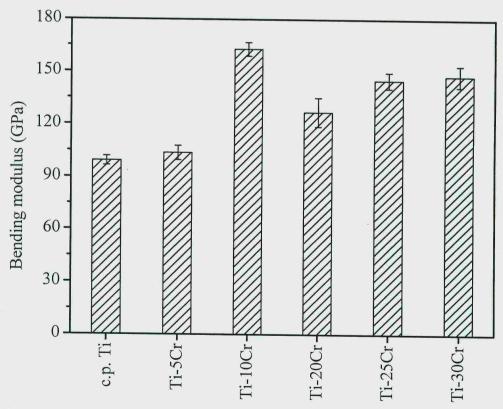


Fig. 2. Bending moduli of c.p. Ti and Ti-Cr alloys.

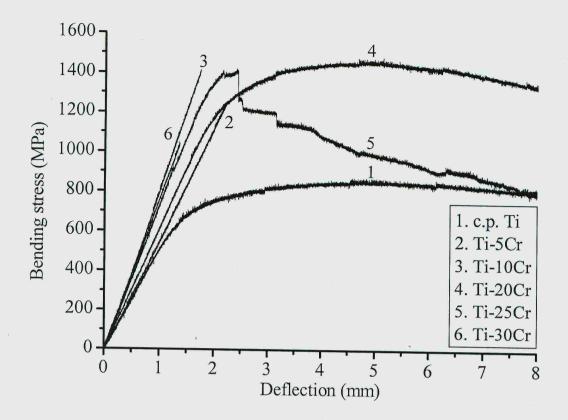


Fig. 3. Bending stress-deflection profiles of c.p. Ti and Ti-Cr alloys.

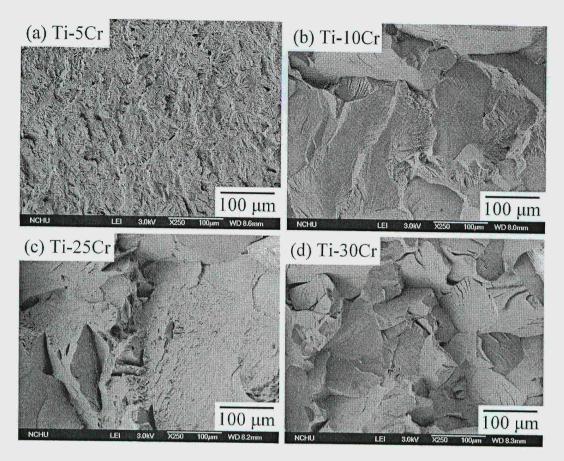


Fig. 4. SEM fractographs of Ti-5Cr (a), Ti-10Cr (b), Ti-25Cr (c) and Ti-30Cr (d) alloys.

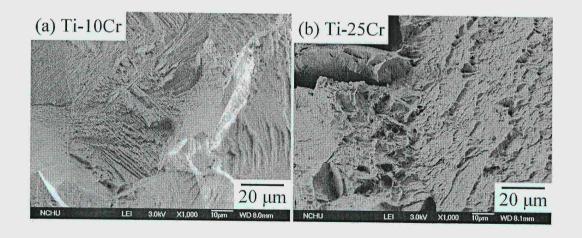


Fig. 5. SEM fractographs of Ti-10Cr (a), Ti-25Cr (b) alloys.

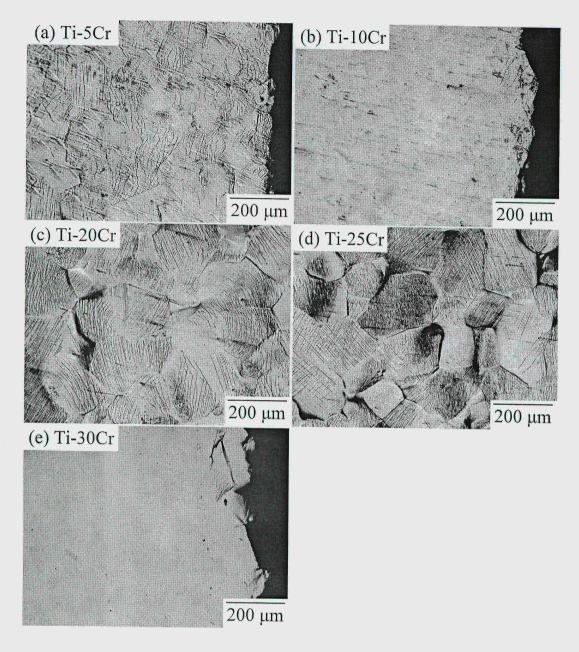


Fig. 6. Optical micrographs of as-bent surfaces of Ti-5Cr (a), Ti-10Cr (b), Ti-20Cr (c), Ti-25Cr (d) and Ti-30Cr (e) alloys.